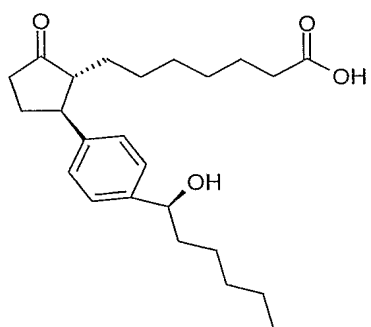


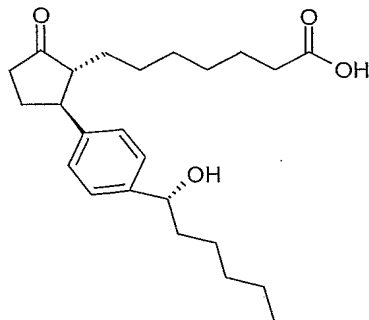
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CLAIMS

1. A compound selected from one of the following:



(1R,2S)-2-[4-(1-(S)-hydroxyhexyl)phenyl]-
5-oxo-cyclopentaneheptanoic acid
[RSS]

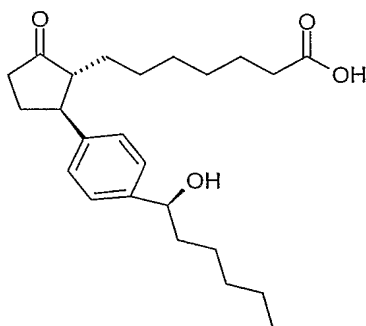


(1R,2S)-2-[4-(1-(R)-hydroxyhexyl)phenyl]-
5-oxo-cyclopentaneheptanoic acid
[RSR]

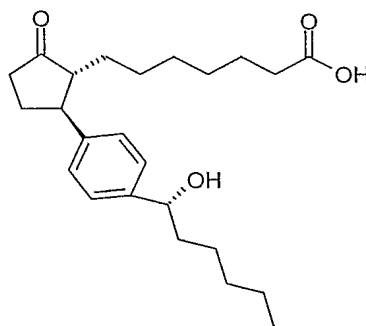
; or ,

5 or a salt, solvate, chemically protected form or prodrug thereof.

2. (*trans*-2-[4-(1-hydroxyhexyl)phenyl]-5-oxo-cyclopentaneheptanoic acid, of which at least 90% by weight
10 is selected from one of the following forms:



(1R,2S)-2-[4-(1-(S)-hydroxyhexyl)phenyl]-
5-oxo-cyclopentaneheptanoic acid
[RSS]



(1R,2S)-2-[4-(1-(R)-hydroxyhexyl)phenyl]-
5-oxo-cyclopentaneheptanoic acid
[RSR]

; or ,

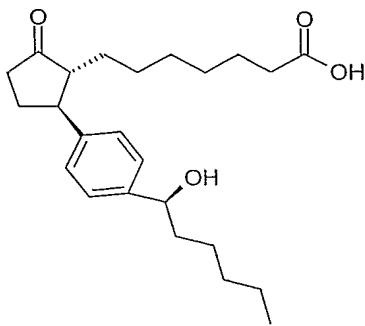
or a salt, solvate, chemically protected form or prodrug thereof.

15

3. 2-[4-(1-hydroxyhexyl)phenyl]-5-oxo-cyclopentaneheptanoic acid, of which at least 80% by weight

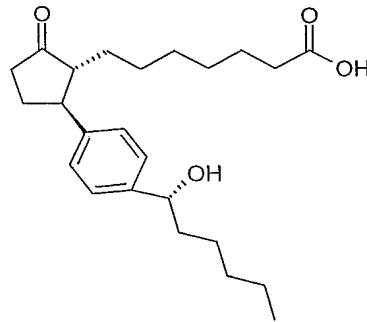
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is in one of the following forms:



(1R,2S)-2-[4-(1-(S)-hydroxyhexyl)phenyl]-
5-oxo-cyclopentaneheptanoic acid
[RSS]

; or



(1R,2S)-2-[4-(1-(R)-hydroxyhexyl)phenyl]-
5-oxo-cyclopentaneheptanoic acid
[RSR]

or a salt, solvate, chemically protected form or prodrug thereof.

5

4. A method of making a compound according to any one of claims 1 to 3.

5. A compound according to any one of claims 1 to 3, or a
10 pharmaceutically acceptable salt thereof, for use in a method of therapy.

6. A pharmaceutical composition comprising a compound
according to any one of claims 1 to 3, or a pharmaceutically
15 acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

7. The use of a compound according to any one of claims 1
to 3, or a pharmaceutically acceptable salt thereof in the
20 preparation of a medicament for the treatment of a condition alleviated by agonism of an EP₂ receptor.

8. The use according to claim 7, wherein the condition
alleviated by agonism of an EP₂ receptor is selected from
25 the group consisting of: glaucoma, dysmenorrhoea and pre-term labour.

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9. A method of treating a condition which can be alleviated by agonism of an EP₂ receptor, which method comprises administering to a patient in need of treatment an effective amount of a compound according to any one of claims 1 to 3, or a pharmaceutically acceptable salt thereof.

10. The method according to claim 9, wherein the condition alleviated by agonism of an EP₂ receptor is selected from the group consisting of: glaucoma, dysmenorrhoea and pre-term labour.

11. The use of an EP₂ receptor agonist, or a pharmaceutically acceptable salt thereof in the preparation of a medicament for the treatment of a condition alleviated by the inhibition of:

- (i) human T-cell activation (proliferation);
- (ii) the release of IL-2;
- (iii) the release of TNF α ; or
- (iv) the release of IFN γ .

12. The use of an EP₂ receptor agonist, or a pharmaceutically acceptable salt thereof in the preparation of a medicament for the treatment of psoriasis.

13. The use of an EP₂ receptor agonist, or a pharmaceutically acceptable salt thereof in the preparation of a medicament for the treatment of inflammatory lung diseases.

14. A use according to any one of claims 11 to 13, wherein the EP₂ receptor agonist is a compound of any one of claims 1 to 3.

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15. A method of treating a condition which can be alleviated by the inhibition of:

- (i) human T-cell activation (proliferation);
- 5 (ii) the release of IL-2;
- (iii) the release of TNF_α ; or
- (iv) the release of IFN_γ ;

which method comprises administering to a patient in need of treatment an effective amount of an EP_2 receptor agonist, or
10 a pharmaceutically acceptable salt thereof.

16. A method of treating a psoriasis, which method comprises administering to a patient in need of treatment an effective amount of an EP_2 receptor agonist, or a
15 pharmaceutically acceptable salt thereof.

17. A method of treating an inflammatory lung disease, which method comprises administering to a patient in need of treatment an effective amount of an EP_2 receptor agonist, or
20 a pharmaceutically acceptable salt thereof.

18. A method according to any one of claims 15 to 17, wherein the EP_2 receptor agonist is a compound of any one of claims 1 to 3.